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C3)

C5)

Amendments To The Claims

1-6. (canceled)

 $7. \label{eq:constraint} \begin{tabular}{ll} T. (previously presented) A compound selected from the following formulae (C1) to (C11) and (C13)-(C20) or a pharmaceutically acceptable salt thereof: \\ \end{tabular}$

C6)

C7)

C8)

C9)

C10)

C11)

C-15)

C16)

C17)

C18)

C19)

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C20)

8. (previously presented) A compound represented by the structural formula AA

or a pharmaceutically acceptable salt thereof.

9. (previously presented) A compound selected from the group consisting of:

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or a pharmaceutically acceptable salt thereof.

10. (canceled)

- (currently amended) A compound according to claim 36 wherein ZFP includes a carboxyl group funcionalized functionalized as a N,N-diethylglycolamido ester, or morpholinylethyl ester.
- 12. (previously presented) The salt derivative of the compound of claim 36 wherein the salt is sodium or potassium.
- 13. (previously presented) A pharmaceutical formulation comprising the compound of claim 36 either with a pharmaceutically acceptable carrier or diluent.

14-17. (canceled)

- 18. (currently amended) A method of treating a mammal to prevent or alleviate the pathological effects of Aone, Actinic keratosis, Insufficient sebum secretion, for Osteoporosis or a maufficient dermal firmness, Insufficient dermal hydration, Psoriasis, Seleroderma, Skin cancer, Skin cell damage from, Mustard vesicants, Wrinkles or Seborrheic dermatitis; wherein the method comprises administering a pharmaceutically effective amount of at least one compound according to claim 36.
 - 19. (original) The method of claim 18 for the treatment of psoriasis.
 - 20. (original) The method of claim 18 for the treatment of osteoporosis.

21-35. (canceled)

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36. (previously presented) A compound represented by a formula below:

wherein

R and R' are independently C₁-C₅ alkyl, or together R and R' form a saturated carbocyclic ring having from 3 to 8 carbon atoms;

RP3 is hydrogen or C1-C5 alkyl;

(L_{P2}) is

where m is 0, 1, or 2,

Zp is a branched C3-C5 alkyl;

ZFB is attached to the 5 or 6 position on the benzofuranyl ring and selected from:

-CO₂H
-CO₂(C₁-C₅ alkyl),
-C(O)NMe₂,
-CO₂(C₁-C₅ alkyl)-NH₂,
-C(O)NH-CH₂-C(O)OH,
-C(O)NH-CH₂-C(O)OEt,
-C(O)NH-CH₂-C(O)OiPr,
-C(O)NH-CH₂-C(O)OBu,
-C(O)NH-CH₂-C(O)OH,
-C(O)NH-CH(Me)-C(O)OH,
-C(O)NH-CH(Me)-C(O)OH,

-C(O)NH-CH(Me)-C(O)OEt, -C(O)NH-CH(Me)-C(O)iPr, -C(O)NH-CH(Me)-C(O)tBu, -C(O)NH-CH(Et)-C(O)OH, -C(O)NH-C(Me)2-C(O)OH, -C(O)NH-C(Me)2-C(O)OMe, -C(O)NH-C(Me)2-C(O)OEt, -C(O)NH-C(Me)2-C(O)iPr. -C(O)NH-C(Me)2-C(O)tBu, -C(O)NH-CMe(Et)-C(O)OH, -C(O)NMe-CH2-C(O)OH, -C(O)NMe-CH2-C(O)OMe, -C(O)NMe-CH2-C(O)OEt, -C(O)NMe-CH2-C(O)OiPr, -C(O)NMe-CH2-C(O)tBu, -C(O)NMe-CH(Me)-C(O)OH, C(O)-NH-5-tetrazolyl, -O-SO2-C1-C5 alkyl), -SO₂(C₁-C₅ alkyl), CH₂S(O)₂Me, CH2S(O)2Et, and CH2S(O)2iPr.

and a pharmaceutically acceptable salt thereof.

37. (previously presented) A compound represented by a formula:

and pharmaceutically acceptable salts thereof.

38. (previously presented) A compound represented by a formula:

and pharmaceutically acceptable salts thereof.

- 39. (withdrawn, currently amended) A method of treating a mammal or alleviating the pathological effects of psoriasis, seleroderma, seborrheic dermatitis or skin cancer or, a mammal in need thereof comprising administered a pharmaceutically effective amount of a compound of Claim 37, or a pharmaceutically acceptable salt thereof.
 - 40. (previously presented) A compound represented by a formula:

and pharmaceutically acceptable salts thereof.

- 41. (previously presented) A pharmaceutical formulation comprising the compound of claim 37 with a pharmaceutically acceptable carrier or diluent.
- (previously presented) A pharmaceutical formulation comprising the compound of claim 38 with a pharmaceutically acceptable carrier or diluent.
- 43. (previously presented) A pharmaceutical formulation comprising the compound of claim 40 with a pharmaceutically acceptable carrier or diluent.